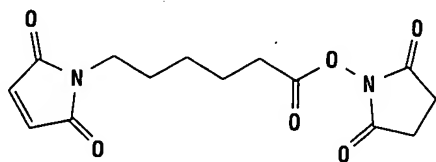


EMCS

Amine-sulphydryl-reactive linker with low immunogenicity and a little more reach.



EMCS
M.W. 308.29
Spacer Arm 9.4 Å

Features/Benefits:

- NHS ester end couples with primary amines at pH 7-9 to form stable amide bonds
- Maleimides react with -SH groups at pH 6.5-7.5, forming stable thioether linkages

- Non-cleavable; water-insoluble
- Increased sphere of coupling vs. GMBS
- Aliphatic spacer offers low potential for eliciting an immune response
- *Reactive groups:* maleimide and NHS ester
- *Reactive toward:* sulphydryl and amino groups
- Literature reference #'s 23, 68, 69 (pages 213-214)

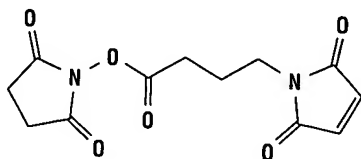
Ordering Information

Product #	Description	Pkg Size	U.S. Price
22308ZZ	EMCS* (N-[ε-Maleimidocaproyloxy]succinimide ester)	50 mg	\$ 70

*Sulfonated, water-soluble analog also available; see Sulfo-EMCS.

GMBS

Heterobifunctional analog of MBS – enhanced maleimide stability with less immunogenicity.



GMBS
M.W. 280.23
Spacer Arm 6.8 Å

Features/Benefits:

- Non-cleavable
- Low potential for eliciting an immune response, ensuring that the primary response to the antigen-carrier protein conjugate is not diluted by a response against a determinant on the cross-linker

- Less immunogenic than SMCC
- *Reactive groups:* NHS ester and maleimide
- *Reactive toward:* amino and sulphydryl groups
- Literature reference #'s 23, 84 (pages 213-214)

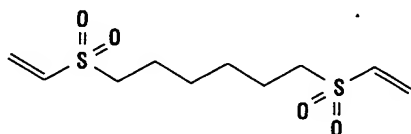
Ordering Information

Product #	Description	Pkg. Size	U.S. Price
22309ZZ	GMBS* (N-[γ-Maleimidobutyryloxy]succinimide ester)	50 mg	\$ 70

*Sulfonated, water-soluble analog also available; see Sulfo-GMBS.

HBVS

Sulphydryl reactivity without the hydrolysis potential of maleimides.



HBVS
M.W. 266.38
Spacer Arm 14.7 Å

Features/Benefits:

- Novel sulphydryl-reactive, homobifunctional cross-linking agent
- Couples via Michael addition, yielding stable thioether links without stereoisomer formation

- Vinylsulfone-reactive group is indefinitely stable at pH 7 and resists hydrolytic degradation for days at pH 9
- Non-cleavable; water-insoluble
- *Reactive group:* vinylsulfone (homobifunctional)
- *Reactive toward:* sulphydryl groups
- Literature reference #'s 78-80 (page 214)

Ordering Information

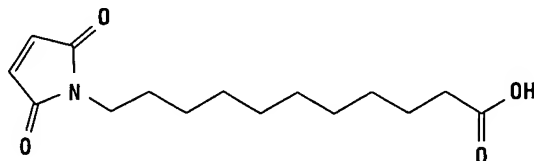
Product #	Description	Pkg. Size	U.S. Price
22344ZZ	HBVS (1,6-Hexane-bis-vinylsulfone)	50 mg	\$ 71

7-3
 Hohenchemie
 3-3
 185

• 800-874-3723

KMUA

Activate biomolecules for cross-linking through sulfhydryl groups or introduce carboxyl groups into proteins.



KMUA
M.W. 281.35
Spacer Arm 15.7 Å

Features/Benefits:

- Novel sulfhydryl-reactive, heterobifunctional cross-linking agent
- Maleimide activate protein/peptide via EDC activation of the carboxyl group
- Sulfhydryl modification agent that creates a terminal carboxylate group at -SH sites in proteins and other molecules

- Maleimide reacts with -SH groups at pH 6.5-7.5, forming stable thioether linkages
- Non-cleavable, long aliphatic cross-bridge
- Useful for preparing peptide-protein conjugates
- *Reactive groups:* maleimide and carboxyl
- *Reactive toward:* sulfhydryl and amino groups
- Literature reference #'s 63, 64 (page 214)

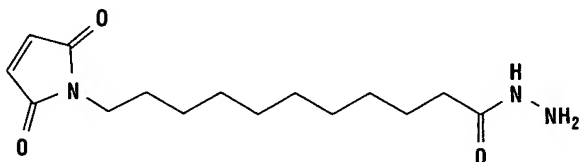
Ordering Information

Product #	Description	Pkg Size	U.S. Price
22211ZZ	KMUA* (N-κ-Maleimidoundecanoic acid)	100 mg	\$ 29

*See also: BMPA, EMCA.

KMUH

Extended chain length; heterobifunctional for the preparation of glycoconjugates.



KMUH
M.W. 295.38
Spacer Arm 19.0 Å

Features/Benefits:

- Sulfhydryl-reactive and carbonyl-reactive heterobifunctional cross-linking agent
- Long, non-cleavable, aliphatic cross-bridge

- Hydrazide group covalently couples to oxidized carbohydrate residues in glycoproteins and other glycoconjugates
- Maleimide reacts with -SH groups at pH 6.5-7.5, forming stable thioether linkages
- *Reactive groups:* maleimide and hydrazide
- *Reactive toward:* sulfhydryl and carbonyl (aldehyde)/carboxyl groups
- Literature reference #67 (page 214)

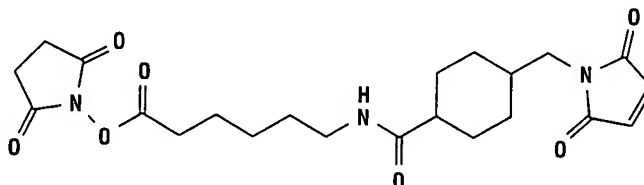
Ordering Information

Product #	Description	Pkg Size	U.S. Price
22111ZZ	KMUH* (N-[κ-Maleimidoundecanoic acid]hydrazide)	50 mg	\$121

*See also: BMPH, EMCH, M,C,H, MPBH, PDPH.

LC-SMCC

Extended chain length analog of SMCC, a popular reagent for immunoconjugate preparation.



LC-SMCC
M.W. 447.48
Spacer Arm 16.1 Å

Features/Benefits:

- Sulfhydryl-reactive and amine-reactive heterobifunctional cross-linking agent
- SMCC and analogs are ideal for coupling enzymes to antibodies as both enzyme activity and antibody specificity can be preserved after coupling
- NHS ester end couples with primary amines at pH 7-9 to form stable amide bonds
- Maleimide reacts with -SH groups at pH 6.5-7.5, forming stable thioether linkages
- Cyclohexane containing cross-bridge stabilizes the maleimide group

- Non-cleavable; water-insoluble
- Useful for the preparation of stable maleimide activated proteins
- *Reactive groups:* NHS ester and maleimide
- *Reactive toward:* amino and sulfhydryl groups
- Literature reference #'s 35, 52, 70 (pages 213-214)

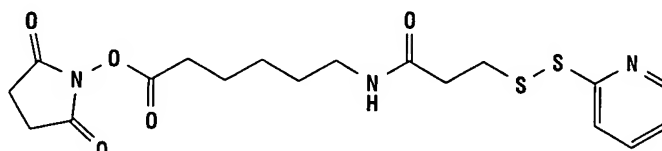
Ordering Information

Product #	Description	Pkg. Size	U.S. Price
22362ZZ	LC-SMCC* (Succinimidyl-4-[N-maleimidomethyl]-cyclohexane-1-carboxy-[6-amidocaproate])	50 mg	\$ 89

*See also: SMCC, Sulfo-SMCC.

LC-SPDP

Classic heterobifunctional, cleavable cross-linker, with an extended spacer arm.



LC-SPDP
M.W. 425.52
Spacer Arm 15.7 Å

Features/Benefits:

- LC-SPDP releases a detectable byproduct after reacting with a free sulfhydryl group; by measuring the release of pyridine-2-thione at 343 nm, the reaction can be easily followed
- *Reactive groups:* pyridyldithio and NHS ester
- *Reactive toward:* sulfhydryl and amino groups
- Literature reference #'s 33, 38 (page 213)

Ordering Information

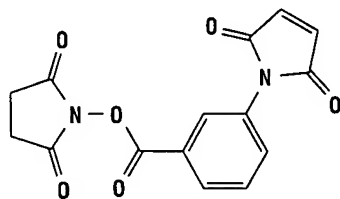
Product #	Description	Pkg. Size	U.S. Price
21651ZZ	LC-SPDP* (Succinimidyl 6-[3-(2-pyridyldithio)-propionamido]hexanoate)	50 mg	\$245

*Sulfonated, water-soluble analog also available; see Sulfo-LC-SPDP.

• 800-874-3723

MBS

Useful for coupling proteins, enzymes to antibodies, toxins to antibodies, and haptens to carrier proteins.



MBS
M.W. 314.25
Spacer Arm 9.9 Å

Features/Benefits:

- Water-insoluble
- Non-cleavable

- Popular for forming enzyme immunoconjugates
- *Reactive groups:* NHS ester and maleimide
- *Reactive toward:* amino and sulfhydryl groups
- Literature reference #'s 25, 42, 84 (pages 213-214)

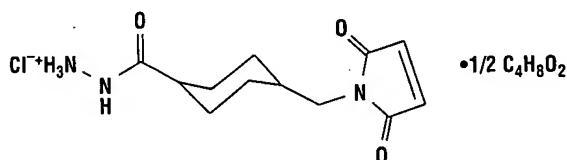
Ordering Information

Product #	Description	Pkg. Size	U.S. Price
22311ZZ	MBS* (<i>m</i> -Maleimidobenzoyl- <i>N</i> -hydroxysuccinimide ester)	50 mg	\$ 41

*Sulfonated, water-soluble analog also available; see Sulfo-MBS.
See also: SMPB.

M₂C₂H

Combine carbohydrate selectivity with sulfhydryl reactivity.



M₂C₂H
M.W. 331.8
Spacer Arm 15.1 Å

Features/Benefits:

- Has an oxidized carbohydrate-specific hydrazide, a sulfhydryl-reactive group and spacer arm to accommodate a wide range of molecular coupling demands
- Sulfhydryl-specific group is a maleimide that yields a thioether linkage upon coupling
- Stable in acetonitrile

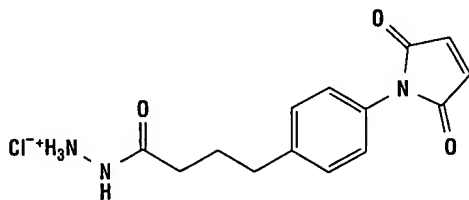
- *Reactive groups:* hydrazide and maleimide
- *Reactive toward:* carbohydrate and sulfhydryl groups
- Literature reference #26 (page 213)

Ordering Information

Product #	Description	Pkg. Size	U.S. Price
22303ZZ	M ₂ C ₂ H (4-[<i>N</i> -Maleimidomethyl]cyclohexane-1-carboxylhydrazide·HCl·1/2 dioxane)	50 mg	\$107

MPBH

Carbohydrate-selective and sulfhydryl-reactive.



MPBH
M.W. 309.75
Spacer Arm 17.9 Å

Features/Benefits:

- Has an oxidized carbohydrate-specific hydrazide, a sulfhydryl-reactive group and spacer arm to accommodate a wide range of molecular coupling demands

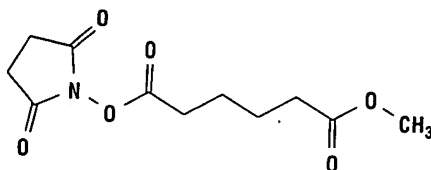
- Sulfhydryl-specific group is a maleimide that yields a thioether linkage upon coupling
- Stable in DMSO
- *Reactive groups:* hydrazide and maleimide
- *Reactive toward:* carbohydrate and sulfhydryl groups
- Literature reference #26 (page 213)

Ordering Information

Product #	Description	Pkg. Size	U.S. Price
22305ZZ	MPBH (4-[4-(<i>N</i> -Maleimidophenyl)]butyric acid hydrazide·HCl)	50 mg	\$104

MSA

Amine-reactive with latent carboxyl group.



MSA
M.W. 257.24
Spacer Arm 7.2 Å

Features/Benefits:

- Amine-reactive modification reagent containing a masked carboxyl group
- NHS ester reacts with primary amines at pH 7-9 to form stable amide bonds

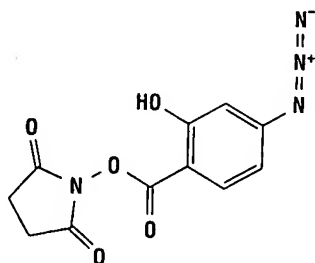
- Masked carboxyl group containing heterobifunctional reagent
- Converts amino groups to carboxyl groups
- Carboxyl group freed at pH 9.5 in phosphate buffer
- Non-cleavable; water-insoluble
- *Reactive groups:* NHS ester and methyl ester
- *Reactive toward:* amino groups

Ordering Information

Product #	Description	Pkg. Size	U.S. Price
22605ZZ	MSA (Methyl <i>N</i> -succinimidyl adipate)	50 mg	\$ 45

NHS-ASA

An ¹²⁵I label can be easily and effectively incorporated into this reagent before the acylation step, and used to radiolabel conjugates.



NHS-ASA
M.W. 276.21
Spacer Arm 8.0 Å

Features/Benefits:

- Photolysis reaction is readily initiated by long wave UV light
- NHS-ASA has been used to simplify detection of photo-affinity labeled complexes and the determination of cross-linked loci
- *Reactive groups:* hydroxyphenyl azide and NHS ester
- *Reactive toward:* amino groups
- Literature reference #27 (page 213)

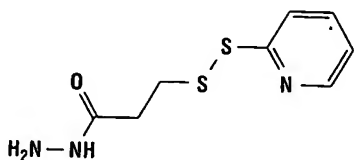
Ordering Information

Product #	Description	Pkg. Size	U.S. Price
27714ZZ	NHS-ASA* (<i>N</i> -Hydroxysuccinimidyl-4-azidosalicylic acid)	50 mg	\$ 65

*Sulfonated, water-soluble analogs also available; see Sulfo-NHS-ASA and Sulfo-LC-NHS-ASA.

PDPH

A cleavable, carbohydrate-selective, sulfhydryl-reactive cross-linker.



PDPH
M.W. 229.32
Spacer Arm 9.2 Å

Features/Benefits:

- Pyridyl disulfide group of PDPH gives a cleavable disulfide linkage in the conjugate

- An oxidized carbohydrate-specific hydrazide
- *Reactive groups:* pyridyldithio and hydrazide
- *Reactive toward:* sulfhydryl groups and carbohydrate
- Literature reference #'s 28, 76, 77 (pages 213-214)

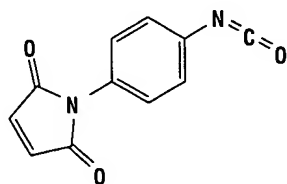
Ordering Information

Product #	Description	Pkg. Size	U.S. Price
22301ZZ	PDPH (3-[2-Pyridyldithio]propionyl hydrazide)	50 mg	\$102

• 800-874-3723

PMPI

Both hydroxyl and sulfhydryl reactivity can be found in this unique cross-linker.



PMPI
M.W. 214.18
Spacer Arm 8.7 Å

Features/Benefits:

- Novel sulfhydryl- and hydroxyl-reactive heterobifunctional cross-linker
- Maleimide reacts with -SH groups at pH 6.5-7.5, forming stable thioether linkages

- Isocyanate reacts with -OH groups to form a carbamate link at pH 8.5
- Non-cleavable
- Excellent tool for the preparation of conjugates of -OH group containing compounds such as steroids and vitamins
- *Reactive groups:* maleimide and isocyanate
- *Reactive toward:* sulfhydryl and hydroxyl groups
- Literature reference #71 (page 214)

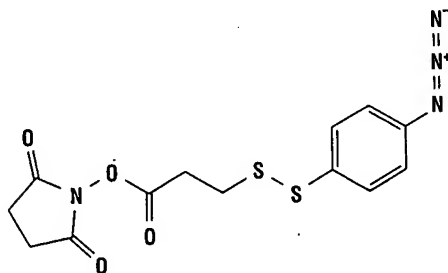
Ordering Information

Product #	Description	Pkg. Size	U.S. Price
28100ZZ	PMPI* (N-[p-Maleimidophenyl]isocyanate)	50 mg	\$ 86

*See also: BMPA, EMCA.

SADP

Versatile, cleavable and photoreactive cross-linker.



SADP
M.W. 352.39
Spacer Arm 13.9 Å

Features/Benefits:

- Cleavable by 50 mM dithiothreitol, 100 mM β-mercaptoethanol or 1% sodium borohydride
- Photolysis is achieved by irradiation at 265-275 nm

- *Reactive groups:* phenylazide and NHS ester
- *Reactive toward:* amino groups
- Literature reference #53 (page 214)

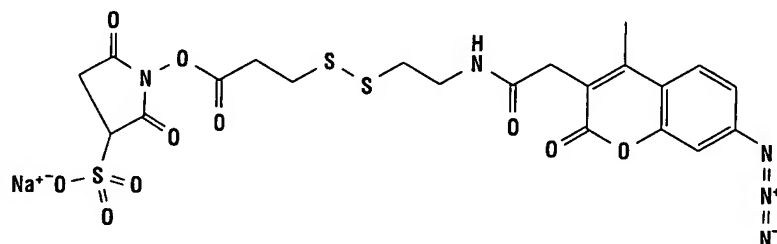
Ordering Information

Product #	Description	Pkg. Size	U.S. Price
21533ZZ	SADP* (N-Succinimidyl [4-azidophenyl]-1,3'-dithiopropionate)	50 mg	\$ 73

*Sulfonated, water-soluble analog also available; see Sulfo-SADP.

SAED

Cleavable and photoreactive – a safer, easier alternative to radiolabeling proteins.



SAED
M.W. 621.60
Spacer Arm 23.6 Å

Features/Benefits:

- *N*-sulfosuccinimidyl ester terminal of SAED will react with amino groups of one protein; the phenyl azide terminal can be reacted nonspecifically under ultraviolet conditions to link a second protein
- Disulfide bond of SAED may be cleaved with an appropriate reducing agent
- *Reactive groups*: azido-methylcoumarin and Sulfo-NHS ester

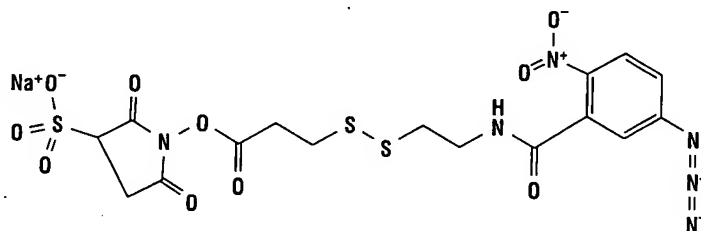
- *Reactive toward*: amino groups
- Literature reference #54 (page 214)

Ordering Information

Product #	Description	Pkg. Size	U.S. Price
33030ZZ	SAED (Sulfosuccinimidyl 2-[7-azido-4-methylcoumarin-3-acetamido]ethyl-1,3'-dithiopropionate)	5 mg	\$103

SAND

Water-soluble, long chain, cleavable analog of ANB-NOS.



SAND
M.W. 570.51
Spacer Arm 18.5 Å

Features/Benefits:

- Cleavable by thiols
- Nitro group on the phenyl azide allows for photolysis at 320-350 nm
- *Reactive groups*: Sulfo-NHS ester and nitrophenyl azide
- *Reactive toward*: amino groups
- Literature reference #29 (page 213)

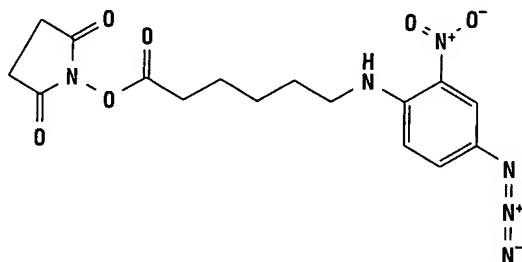
Ordering Information

Product #	Description	Pkg. Size	U.S. Price
21549ZZ	SAND (Sulfosuccinimidyl 2-[<i>m</i> -azido- <i>o</i> -nitrobenzamido]ethyl-1,3'-dithiopropionate)	50 mg	\$131

• 800-874-3723

SANPAH

Extended chain length, photoactivatable cross-linker.



SANPAH
M.W. 390.35
Spacer Arm 18.2 Å

Features/Benefits:

- Optimal photolysis occurs at 320-350 nm; a condition that limits damage to biomolecules by irradiation

- *Reactive groups:* nitrophenyl azide and NHS ester
- *Reactive toward:* amino groups
- Literature reference #30 (page 213)

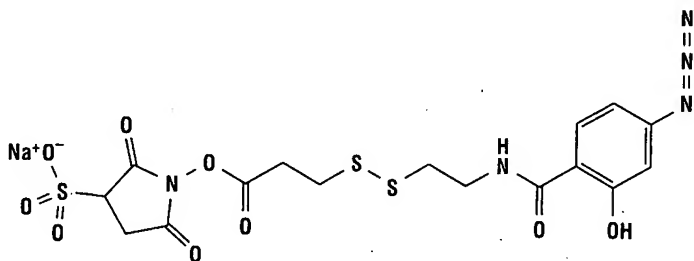
Ordering Information

Product #	Description	Pkg. Size	U. Price
22600ZZ	SANPAH* (N-Succinimidyl 6-[4'-azido-2'-nitro-phenylamino]hexanoate)	50 mg	\$

*Sulfonated, water-soluble analog also available; see Sulfo-SANPAH.

SASD

Transfers a radioactive label from one protein to another.



SASD
M.W. 541.51
Spacer Arm 18.9 Å

Features/Benefits:

- Radioiodinatable,* cleavable, photoreactive, heterobifunctional cross-linker
- Iodination occurs between the azide and the hydroxyl groups of the phenyl ring; after cleavage by a reducing agent, the radioactive label will remain attached to the protein conjugated by photolysis

- *Reactive groups:* hydroxyphenyl azide and Sulfo-NHS ester
- *Reactive toward:* amino groups
- Literature reference #'s 31, 51 (pages 213-214)

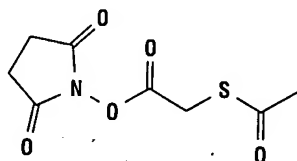
Ordering Information

Product #	Description	Pkg. Size	U. Price
27716ZZ	SASD (Sulfosuccinimidyl-2-[p-azido-salicylamido]ethyl-1,3'-dithiopropionate)	50 mg	\$1

See IODO-GEN Iodination Reagent (Product #'s 28600ZZ and 28601ZZ).

SATA

Adds protected sulfhydryls for better control when a sulfhydryl group is needed in conjugate formation.



SATA
M.W. 231.23
Spacer Arm 2.8 Å

Features/Benefits:

- Reacts with amines to add protected sulfhydryl groups
- NHS ester end couples with primary amines at pH 7-9 to form stable amide bonds
- Converts amino groups to sulfhydryl groups
- Latent -SH group is released by hydroxylamine and is available for reaction with maleimide-activated biomolecules or other -SH group-containing compounds

- Cross-links formed with other -SH group-containing molecules are reversible by reducing agents
- *Reactive groups:* NHS ester and thioacetyl-protected sulfhydryl
- *Reactive toward:* amino and maleimide/iodoacetyl or vinyl sulfone
- Literature reference #72 (page 214)

Ordering Information

Product #	Description	Pkg Size	U.S. Price
26102ZZ	SATA* (N-Succinimidyl S-acetylthioacetate)	50 mg	\$ 30

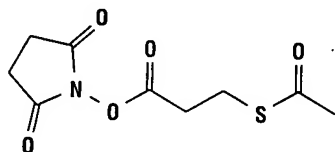
*See also: SATP.

Compatible Pierce Products for Use With SATA and SATP

20688ZZ	DMSO	950 ml	\$ 29
26103ZZ	Hydroxylamine•HCl	25 gm	\$ 28

SATP

Same function as SATA, but offers more steric freedom for the unmasked sulfhydryl group.



SATP
M.W. 245.25
Spacer Arm 4.1 Å

Features/Benefits:

- Thiolation reagent that reacts with amines to add protected sulfhydryl groups
- Converts amino groups to sulfhydryl groups
- NHS ester end couples with primary amines at pH 7-9 to form stable amide bonds
- Latent -SH group is released by hydroxylamine treatment and available for reaction with maleimide-activated biomolecules or other -SH group-containing compounds

- Cross-links formed with other -SH group-containing molecules are reversible by reducing agents
- Modify peptides to facilitate the preparation of hapten-carrier conjugates
- *Reactive groups:* NHS ester and thioacetyl-protected sulfhydryl
- *Reactive toward:* amino and maleimide/iodoacetyl or vinyl sulfone
- Literature reference #'s 72, 81 (page 214)

Ordering Information

Product #	Description	Pkg Size	U.S. Price
26100ZZ	SATP (N-Succinimidyl S-acetylthiopropionate)	50 mg	\$ 49

Compatible Pierce Products for Use With SATP

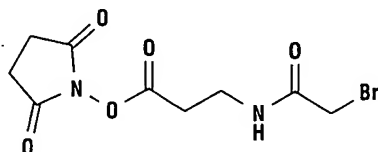
20688ZZ	DMSO	950 ml	\$ 29
26103ZZ	Hydroxylamine•HCl	25 gm	\$ 28

Pierce Chemical Company

• 800-874-3723

SBAP

Alternative active halogen chemistry applied to the preparation of cyclic peptides and peptide conjugates.



SBAP
M.W. 307.10
Spacer Arm 6.2 Å

Features/Benefits:

- NHS ester reacts with primary amines at pH 7-9 to form a stable amide bond
- Bromoacetyl group reacts with sulfhydryl groups at pH > 7.5 to form stable thioether bonds

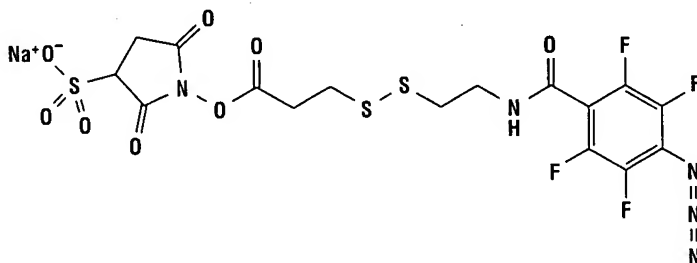
- Spacer maintains peptide-like character in the cross-linked species
- Resulting cross-link is susceptible to acid hydrolysis
- *Reactive groups:* NHS ester and bromoacetyl
- *Reactive toward:* amino and sulfhydryl groups
- Literature reference #73 (page 214)

Ordering Information

Product #	Description	Pkg Size	U.S. Price
22339ZZ	SBAP (Succinimidyl 3-[bromoacetamido]propionate)	50 mg	\$ 72

SFAD

Significantly improved photoconjugation efficiency over typical aryl azide-containing cross-linkers.



SFAD
M.W. 597.48
Spacer Arm 14.6 Å

Features/Benefits:

- Water-soluble; cleavable
- Perfluorophenyl azide moiety photolyzes at 320 nm
- Insertion efficiency approximately 70%
- Improved stability of the singlet perfluoroaryl nitrene reactive intermediate allows high-efficiency insertion with -CH bonds vs. low efficiency ring expansion with amine nucleophiles, typical of nonfluorinated aryl nitrenes
- ¹⁹F NMR can be used to monitor perfluoroaryl moiety transfer from one protein to another

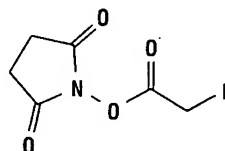
- *Reactive groups:* Sulfo-NHS ester and perfluoroaryl azide moiety
- *Reactive toward:* amino groups and -CH bonds
- Literature reference #'s 82, 83 (page 214)

Ordering Information

Product #	Description	Pkg. Size	U.S. Price
27719ZZ	SFAD (Sulfosuccinimidyl-[perfluoroazidobenzamido]-ethyl-1,3'-dithiopropionate)	50 mg	\$109

SIA

Suited to capturing amine and sulphydryl groups in very close proximity.



SIA
M.W. 283.02
Spacer Arm 1.5 Å

Features/Benefits:

- Shortest sulphydryl-reactive and amine-reactive heterobifunctional cross-linker available
- Non-cleavable; close proximity cross-linking agent

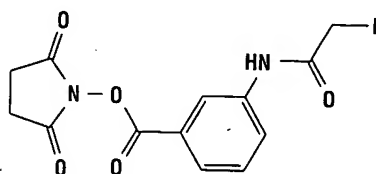
- NHS ester reacts with primary amines at pH 7-9 to form a stable amide bond
- Iodoacetyl group reacts with sulphydryl groups at pH > 7.5 to form stable thioether bond
- *Reactive groups:* NHS ester and iodoacetyl
- *Reactive toward:* amino and sulphydryl groups
- Literature reference #'s 74, 75 (page 214)

Ordering Information

Product #	Description	Pkg. Size	U.S. Price
22349ZZ	SIA (N-Succinimidyl iodoacetate)	50 mg	\$ 39

SIAB

Popular alternative for making enzyme-antibody conjugates; reactive toward amines and sulphydryls.



SIAB
M.W. 402.14
Spacer Arm 10.6 Å

Features/Benefits:

- Used to prepare stable enzyme-IgG conjugates
- Forms conjugates with liposomes

- *Reactive groups:* iodoacetyl and NHS ester
- *Reactive toward:* sulphydryl and amino groups
- Literature reference #'s 33, 34 (page 213)

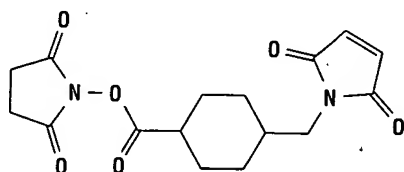
Ordering Information

Product #	Description	Pkg. Size	U.S. Price
22329ZZ	SIAB* (N-Succinimidyl[4-iodoacetyl]aminobenzoate)	50 mg	\$ 89

*Sulfonated, water-soluble analog also available; see Sulfo-SIAB.
See also: SIA, SBAP.

SMCC

Provides stable activated proteins.



SMCC
M.W. 334.32
Spacer Arm 11.6 Å

Features/Benefits:

- Cyclohexane bridge gives extra stability to the maleimide-reactive group; N-(4-carboxycyclohexylmethyl) maleimide groups are stable for 64 hours (in 0.1 M sodium phosphate buffer, pH 7.0 at 4°C)

- Ideal for coupling enzymes to antibodies; both the enzyme activity and antibody specificity can be preserved after coupling
- *Reactive groups:* NHS ester and maleimide
- *Reactive toward:* amino and sulphydryl groups
- Literature reference #'s 35, 52, 84, 89 (pages 213-214)

Ordering Information

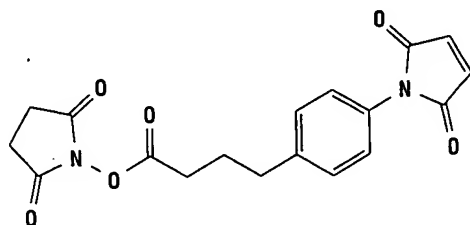
Product #	Description	Pkg. Size	U.S. Price
22360ZZ	SMCC* (Succinimidyl 4-[N-maleimidomethyl]-cyclohexane-1-carboxylate)	50 mg	\$ 54

*Sulfonated, water-soluble analog also available; see Sulfo-SMCC.
See also: AMAS, BMPS, MBS, SMPB, SMPH.

• 800-874-3723

SMPB

Features extended chain length to limit steric hindrance.



SMPB
M.W. 356.33
Spacer Arm 11.6 Å

Features/Benefits:

- Extended chain analog of MBS
- Conjugates formed with SMPB were shown to be more stable in serum than SPDP conjugates

- *Reactive groups:* NHS ester and maleimide
- *Reactive toward:* amino and sulfhydryl groups
- Literature reference #'s 36, 84 (pages 213-214)

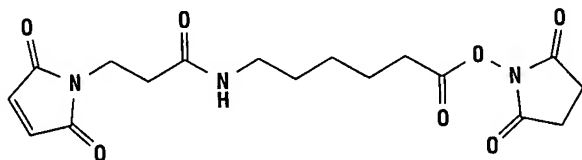
Ordering Information

Product #	Description	Pkg. Size	U.S. Price
22416ZZ	SMPB* (Succinimidyl 4-[p-maleimidophenyl]-butyrate)	50 mg	\$ 65

*Sulfonated, water-soluble analog also available; see Sulfo-SMPB.

SMPH

Spacer arm longer than that of SMPB, minimizing steric hindrance.



SMPH
M.W. 379.36
Spacer Arm 14.3 Å

Features/Benefits:

- Amine- and sulfhydryl-reactive heterobifunctional cross-linker
- NHS ester reacts with primary amines at pH 7-9 to form a stable amide bond

- Maleimide reacts with -SH groups at a pH of 6.5-7.5, forming stable thioether bond
- Non-cleavable; water-insoluble
- More hydrophilic character to the cross-bridge than BMPS, GMBS, EMCS or SMPB
- *Reactive groups:* maleimide and NHS ester
- *Reactive toward:* sulfhydryl and amino groups
- Literature reference #'s 74, 75 (page 214)

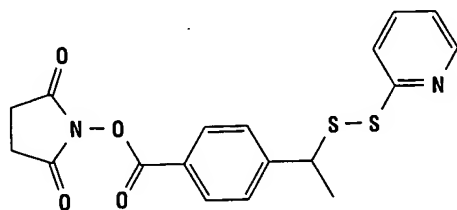
Ordering Information

Product #	Description	Pkg. Size	U.S. Price
22363ZZ	SMPH* (Succinimidyl-6-[(β-maleimidopropionamido)hexanoate])	50 mg	\$ 89

*See also: BMPS, GMBS, EMCS, SMPB.

SMPT

Forms cleavable immunotoxins with greater stability in vivo.



SMPT
M.W. 388.46
Spacer Arm 20.0 Å

Features/Benefits:

- Contains a hindered disulfide bond; has formed immunotoxins with improved stability

- *In vitro*, an SMPT conjugate was as effective as conjugates formed with SPDP and 2-Iminothiolane
- Does not require exposing the antibody to reducing agents
- *Reactive groups:* NHS ester and pyridyldithio
- *Reactive toward:* amino and sulfhydryl groups
- Literature reference #37 (page 213)

Ordering Information

Product #	Description	Pkg. Size	U.S. Price
21558ZZ	SMPT* (4-Succinimidylloxycarbonyl-methyl-α-[2-pyridyldithio]toluene)	50 mg	\$161

*Sulfonated, long chain water-soluble analog also available; see Sulfo-LC-SMPT.

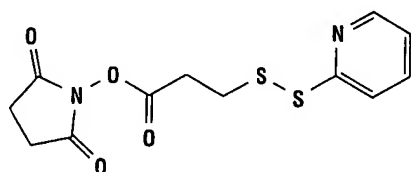
SPB

Amine-reactive DNA intercalating agent that has applications in cross-linking protein to DNA and photoimmobilizing compounds onto microwell plates.

See Product # 23013ZZ, (Succinimidyl-[4-(psoralen-8-yloxy)]butyrate), in Section 3-4, page 224.

SPDP

Classic heterobifunctional, cleavable cross-linker.



SPDP
M.W. 312.37
Spacer Arm 6.8 Å

Features/Benefits:

- Widely used in immunochemistry; conjugates used in drug carrier systems, antibody production and enzyme immunoassays have been successfully prepared with SPDP

- SPDP can be used as a protein thiolation reagent, resulting in available -SH groups
- *Reactive groups:* NHS ester and pyridyldithio
- *Reactive toward:* amino and sulfhydryl groups
- Literature reference #'s 38, 86 (pages 213-214)

Ordering Information

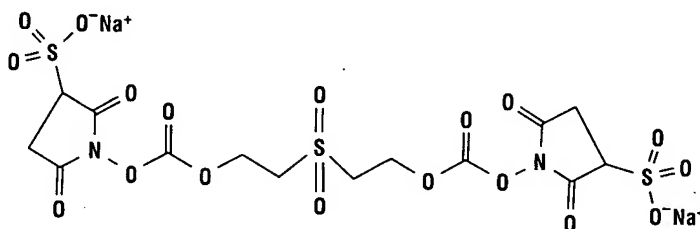
Product #	Description	Pkg. Size	U.S. Price
21857ZZ	SPDP* (N-Succinimidyl 3-[2-pyridyldithio]propionate)	50 mg	\$ 85

*See: LC-SPDP, Sulfo-LC-SPDP.

See also: SATA, SATP (protein thiolation reagents).

Sulfo-BSOCOES

Water-soluble, base-reversible analog of BSOCOES.



Sulfo-BSOCOES
M.W. 640.44
Spacer Arm 13.0 Å

Features/Benefits:

- Water-soluble
- Base-cleavable (pH 11.6, 2 hours, 37°C)
- *Reactive groups:* Sulfo-NHS ester (homobifunctional)
- *Reactive toward:* amino groups
- Literature reference #39 (page 213)

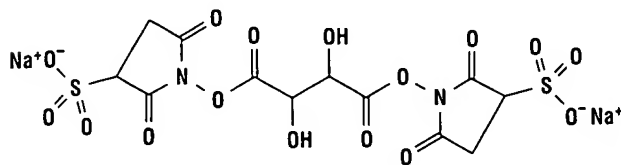
Ordering Information

Product #	Description	Pkg. Size	U.S. Price
21556ZZ	Sulfo-BSOCOES (Bis[2-(Sulfosuccinimidooxycarbonyloxy)-ethyl]sulfone)	50 mg	\$103

• 800-874-3723

Sulfo-DST

Water-soluble analog of DST; cleavable by oxidizing agents.



Sulfo-DST
M.W. 548.32
Spacer Arm 6.4 Å

Features/Benefits:

- Ideal for applications in which a cross-link reversibility is desired without disturbing protein S-S bonds

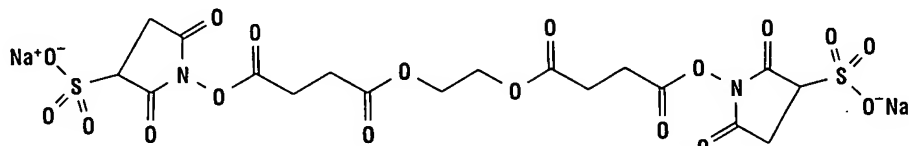
- *Reactive groups:* Sulfo-NHS ester (homobifunctional)
- *Reactive toward:* amino groups
- Literature reference #40 (page 213)

Ordering Information

Product #	Description	Pkg. Size	U.S. Price
20591ZZ	Sulfo-DST (Disulfosuccinimidyl tartrate)	50 mg	\$ 98

Sulfo-EGS

Water-soluble and cleavable – without harsh reducing agents.



Sulfo-EGS
M.W. 660.45
Spacer Arm 16.1 Å

Features/Benefits:

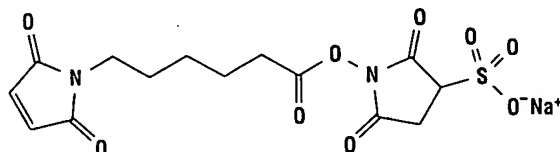
- Cross-links formed are cleavable at pH 8.5 using hydroxylamine for three to six hours at 37°C
- Lactose dehydrogenase retained 60% of its activity after reversible cross-linking with EGS
- *Reactive groups:* Sulfo-NHS esters (homobifunctional)
- *Reactive toward:* amino groups
- Literature reference #41 (page 213)

Ordering Information

Product #	Description	Pkg. Size	U.S. Price
21566ZZ	Sulfo-EGS (Ethylene glycol bis[sulfosuccinimidylsuccinate])	50 mg	\$ 95

Sulfo-EMCS

Water-soluble, low immunogenicity and a little more reach.



Sulfo-EMCS
M.W. 410.33
Spacer Arm 9.4 Å

Features/Benefits:

- Sulfonated analog of EMCS has improved solubility in aqueous buffer systems
- Sulfo-NHS ester end couples with primary amines at pH 7-9 to form stable amide bonds
- Maleimide reacts with -SH groups at pH 6.5-7.5, forming stable thioether linkages

- Non-cleavable
- Increased sphere of coupling vs. Sulfo-GMBS
- Aliphatic spacer offers low potential for eliciting an immune response
- *Reactive groups:* maleimide and Sulfo-NHS ester
- *Reactive toward:* sulfhydryl and amino groups
- Literature reference #'s 23, 68, 69 (pages 213-214)

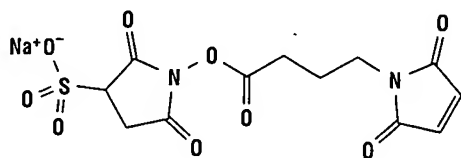
Ordering Information

Product #	Description	Pkg. Size	U.S. Price
22307ZZ	Sulfo-EMCS* (N-[ε-Maleimidocaproyloxy] sulfosuccinimide ester)	50 mg	\$149

*Non-sulfonated, water-insoluble analog also available; see EMCS.
See also: Sulfo-GMBS and Sulfo-KMUS.

Sulfo-GMBS

Sulfonated analog of GMBS offers water solubility, enhanced stability with less immunogenicity.



Sulfo-GMBS
M.W. 382.28
Spacer Arm 6.8 Å

Features/Benefits:

- Non-cleavable, membrane-impermeable
- Has a low potential for eliciting an immune response, ensuring that the primary response to the antigen-carrier protein conjugate is not diluted by a response against a determinant on the cross-linker

- GMBS and Sulfo-GMBS are reported to be less immunogenic than SMCC
- Sulfo-GMBS has improved solubility in water and aqueous buffer systems
- *Reactive groups:* maleimide and Sulfo-NHS ester
- *Reactive toward:* sulfhydryl and amino groups
- Literature reference #23 (page 213)

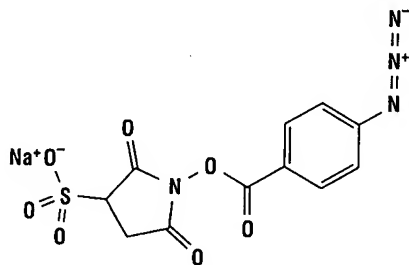
Ordering Information

Product #	Description	Pkg. Size	U.S. Price
22324ZZ	Sulfo-GMBS* (N-[γ-Maleimidobutyryloxy]sulfo-succinimide ester)	50 mg	\$137

*See also: Sulfo-EMCS, Sulfo-KMUS, Sulfo-MBS.

Sulfo-HSAB

Water-soluble, amine-reactive, photoreactive cross-linker.



Sulfo-HSAB
M.W. 362.25
Spacer Arm 9.0 Å

Features/Benefits:

- Useful for the preparation of conjugates for the purpose of producing antibodies

- *Reactive groups:* phenyl azide and Sulfo-NHS ester
- *Reactive toward:* amino groups
- Literature reference #'s 24, 30 (page 213)

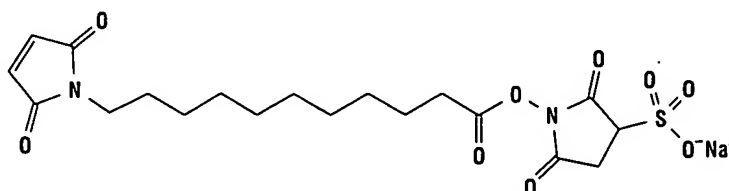
Ordering Information

Product #	Description	Pkg. Size	U.S. Price
21563ZZ	Sulfo-HSAB (N-Hydroxysulfosuccinimidyl-4-azidobenzoate)	50 mg	\$ 54

• 800-874-3723

Sulfo-KMUS

Water-soluble, long chain heterobifunctional cross-linker.



Sulfo-KMUS
M.W. 480.47
Spacer Arm 15.7 Å

Features/Benefits:

- Non-cleavable extended aliphatic cross-bridge
- Water-soluble with enhanced sphere of coupling
- Sulfo-NHS ester reacts with primary amines at pH 7-9 to form a stable amide bond
- Maleimide reacts with -SH groups at pH 6.5-7.5, forming a stable thioether bond
- *Reactive groups:* maleimide and Sulfo-NHS ester
- *Reactive toward:* sulfhydryl and amino groups
- Literature reference #23 (page 213)

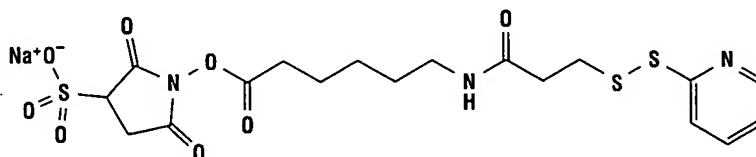
Ordering Information

Product #	Description	Pkg Size	U.S. Price
21111ZZ	Sulfo-KMUS* (N-[α -Maleimidoundecanoyloxy]-sulfosuccinimide ester)	50 mg	\$159

*See also: Sulfo-GMBS, Sulfo-EMCS.

Sulfo-LC-SPDP

Classic water-soluble, heterobifunctional, cleavable cross-linker.



Sulfo-LC-SPDP
M.W. 527.57
Spacer Arm 15.6 Å

Features/Benefits:

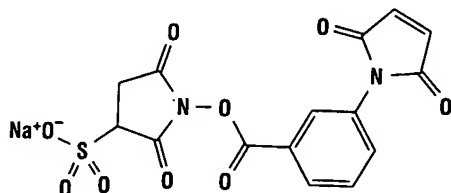
- Sulfo-LC-SPDP releases a detectable byproduct after reacting with free sulfhydryl groups; by measuring the release of pyridine-2-thione at 343 nm, the reaction can be easily followed
- *Reactive groups:* pyridyldithio and Sulfo-NHS ester
- *Reactive toward:* sulfhydryl and amino groups
- Literature reference #'s 38, 85 (pages 213-214)

Ordering Information

Product #	Description	Pkg. Size	U.S. Price
21650ZZ	Sulfo-LC-SPDP (Sulfosuccinimidyl 6-[3'-(2-pyridyldithio)-propionamido]hexanoate)	50 mg	\$257

Sulfo-MBS

Useful for coupling proteins, enzymes to antibodies, toxins to antibodies, and haptens to carrier proteins.



Sulfo-MBS
M.W. 416.30
Spacer Arm 9.9 Å

Features/Benefits:

- Water-soluble; non-cleavable
- Membrane-impermeable

- **Reactive groups:** Sulfo-NHS ester and maleimide
- **Reactive toward:** amino and sulfhydryl groups
- Literature reference #42 (page 213)

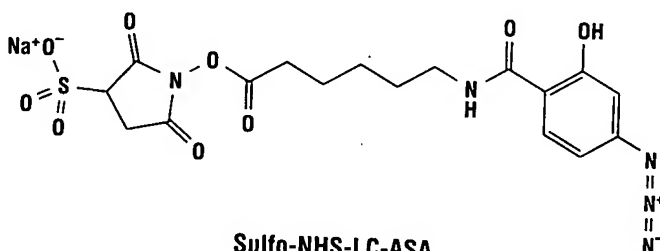
Ordering Information

Product #	Description	Pkg. Size	U.S. Price
22312ZZ	Sulfo-MBS* (<i>m</i> -Maleimidobenzoyl- <i>N</i> -hydroxysulfo-succinimide ester)	50 mg	\$ 94

*See also: Sulfo-EMCS, Sulfo-GMBS, Sulfo-KMUS.

Sulfo-NHS-LC-ASA

An ¹²⁵I label can be easily and effectively incorporated into this reagent before the acylation step and used to radiolabel conjugates.



Sulfo-NHS-LC-ASA
M.W. 491.41
Spacer Arm 18.0 Å

Features/Benefits:

- Photolysis reaction is readily initiated by long wave UV light
- Simplifies detection of photoaffinity labeled complexes and the determination of cross-linked loci
- Sulfo-NHS-LC-ASA is water-soluble, and has a long spacer arm to overcome steric restraints
- **Reactive groups:** hydroxy phenyl azide and Sulfo-NHS ester
- **Reactive toward:** amino groups
- Literature reference #27 (page 213)

Ordering Information

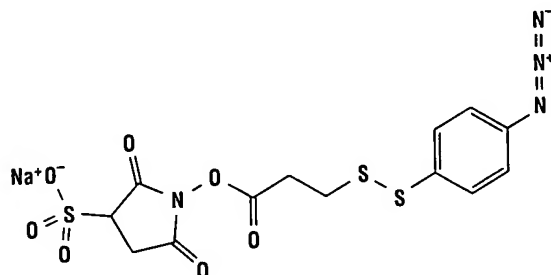
Product #	Description	Pkg. Size	U.S. Price
27735ZZ	Sulfo-NHS-LC-ASA* (Sulfosuccinimidyl[4-azidosalicylamido]-hexanoate)	50 mg	\$225

*See also: SASD.

• 800-874-3723

Sulfo-SADP

Highly versatile, water-soluble, cleavable and photoreactive cross-linker.



Sulfo-SADP
M.W. 454.44
Spacer Arm 13.9 Å

Features/Benefits:

- Cleavable by 50 mM dithiothreitol, 100 mM β-mercaptoethanol or 1% sodium borohydride

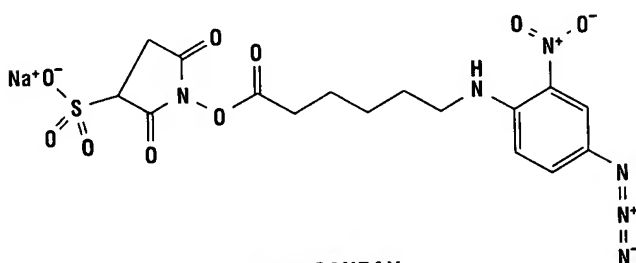
- Photolysis is achieved by irradiation at 265-275 nm
- *Reactive groups:* phenyl azide and Sulfo-NHS ester
- *Reactive toward:* amino groups
- Literature reference #20 (page 213)

Ordering Information

Product #	Description	Pkg. Size	U.S. Price
21553ZZ	Sulfo-SADP (Sulfosuccinimidyl[4-azidophenylthio]-propionate)	50 mg	\$125

Sulfo-SANPAH

Extended chain length, photoactivatable cross-linker.



Sulfo-SANPAH
M.W. 492.40
Spacer Arm 18.2 Å

Features/Benefits:

- Optimal photolysis occurs at 320-350 nm, a condition that limits damage to biomolecules by irradiation

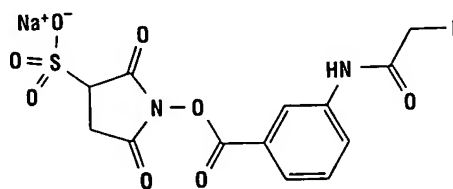
- Water-soluble; non-cleavable
- *Reactive groups:* nitrophenyl azide and Sulfo-NHS ester
- *Reactive toward:* amino groups
- Literature reference #30 (page 213)

Ordering Information

Product #	Description	Pkg. Size	U.S. Price
22589ZZ	Sulfo-SANPAH (Sulfosuccinimidyl 6-[4'-azido-2'-nitro-phenylamino]hexanoate)	50 mg	\$139

Sulfo-SIAB

Popular alternative for making enzyme-antibody conjugates - reactive toward amines and sulphydryls.



Sulfo-SIAB
M.W. 504.19
Spacer Arm 10.6 Å

Features/Benefits:

- *Reactive groups:* iodoacetate and Sulfo-NHS ester
- *Reactive toward:* amino and sulphydryl groups
- Literature reference #45 (page 214)

Ordering Information

Product #	Description	Pkg. Size	U.S. Price
22327ZZ	Sulfo-SIAB (Sulfosuccinimidyl[4-iodoacetyl]aminobenzoate)	50 mg	\$ 99

Special Product

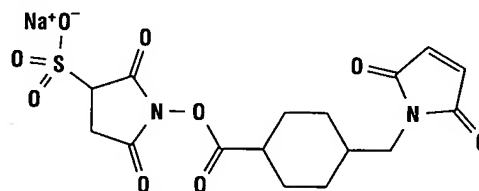


Sulfo-SMCC

Provides stable activated proteins.

Features/Benefits:

- Water-soluble, non-cleavable and membrane-impermeable
- Cyclohexane bridge gives extra stability to the maleimide-reactive group; *N*-(4-carboxycyclohexylmethyl)maleimide groups are stable for 64 hours (in 0.1 M sodium phosphate buffer, pH 7.0 at 4°C)
- Ideal for coupling enzymes to antibodies; both the enzyme activity and antibody specificity can be preserved after coupling
- *Reactive groups:* Sulfo-NHS ester and maleimide
- *Reactive toward:* amino and sulfhydryl groups
- Literature reference #46 (page 214)



Sulfo-SMCC
M.W. 436.37
Spacer Arm 11.6 Å

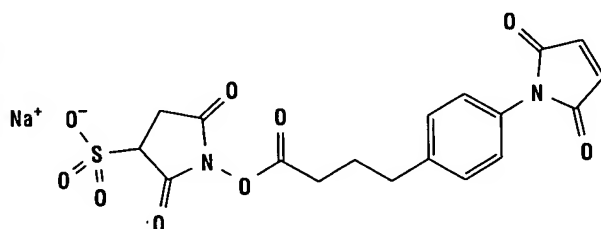
Ordering Information

Product #	Description	Pkg. Size	U.S. Price
22322ZZ	Sulfo-SMCC (Sulfosuccinimidyl 4-[<i>N</i> -maleimidomethyl]-cyclohexane-1-carboxylate)	50 mg	\$104

• 800-874-3723

Sulfo-SMPB

Extended chain length analog of Sulfo-MBS.



Sulfo-SMPB
M.W. 458.38
Spacer Arm 14.5 Å

Features/Benefits:

- Extended chain length limits steric hindrance
- Water-soluble; non-cleavable

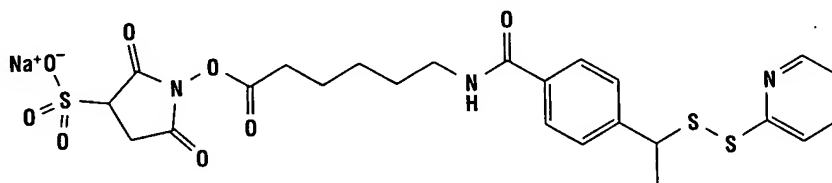
- Membrane-impermeable
- *Reactive groups:* Sulfo-NHS ester and maleimide
- *Reactive toward:* amino and sulfhydryl groups
- Literature reference #'s 36, 47 (pages 213-214)

Ordering Information

Product #	Description	Pkg. Size	U.S. Price
22317ZZ	Sulfo-SMPB (Sulfosuccinimidyl 4-[p-maleimidophenyl]-butyrate)	50 mg	\$101

Sulfo-LC-SMPT

Form cleavable immunotoxins with greater stability in vivo.



Sulfo-LC-SMPT
M.W. 603.67
Spacer Arm 20.0 Å

Features/Benefits:

- Contains a hindered disulfide bond; has formed immunotoxins with improved stability
- *In vitro*, SMPT conjugates are as effective as conjugates formed with SPDP and 2-Iminoethanol
- Does not require exposing the antibody to reducing agents
- Offers an extended spacer arm and water solubility

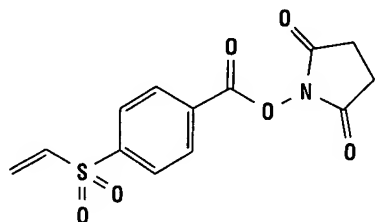
- *Reactive groups:* Sulfo-NHS ester and pyridyldithio
- *Reactive toward:* amino and sulfhydryl groups
- Literature reference #48 (page 214)

Ordering Information

Product #	Description	Pkg. Size	U.S. Price
21568ZZ	Sulfo-LC-SMPT (Sulfosuccinimidyl-6-[α-methyl-α-(2-pyridyldithio)toluamido]hexanoate)	50 mg	\$229

SVSB

Combines the advantages of the vinylsulfone-reactive group with those of the classical NHS ester.



SVSB
M.W. 309.30
Spacer Arm 8.3 Å

Features/Benefits:

- Novel heterobifunctional reagent containing an amine-reactive NHS and a sulfhydryl-reactive vinylsulfone
- Non-cleavable; water-insoluble
- NHS ester reacts with primary amines at pH 7-9 to form a stable amide bond

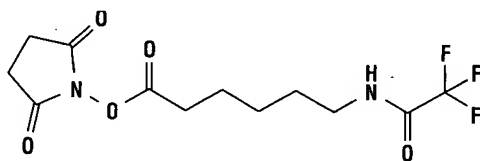
- Vinylsulfone reacts with -SH groups, forming a stable thioether linkage
- Conjugates prepared with vinylsulfone-reactive group form a single stereoisomer
- Unlike the maleimide active group, vinylsulfone is not subject to hydrolytic degradation in aqueous environments
- After modification of an amine-containing molecule with SVSB, the vinylsulfone intermediate can be stored for long periods without loss of sulfhydryl reactivity
- *Reactive groups:* vinylsulfone and NHS ester
- *Reactive toward:* sulfhydryl and amino groups
- Literature reference #'s 78-80 (page 214)

Ordering Information

Product #	Description	Pkg Size	U.S. Price
22358ZZ	SVSB (N-Succinimidyl-[4-vinylsulfonyl]benzoate)	50 mg	\$107

TFCS

Amine-reactive cross-linker with a latent amino group available on demand.



TFCS
M.W. 324.25
Spacer Arm 7.7 Å

Features/Benefits:

- Amine-reactive modification agent with a protected primary amine group
- NHS ester end couples with primary amines at pH 7-9 to form stable amide bonds
- Used to extend lysine side-chain length to reduce steric hindrance

- Temporarily block amine groups in target molecules
- Trifluoroacetyl protecting group released by phosphate or borate buffer, pH 7.8-8.1
- Unmasked amino group ready for reaction with any amine-reactive cross-linker for conjugate preparation
- Non-cleavable; water-insoluble
- *Reactive groups:* NHS ester and trifluoroacetyl-protected -NH₂
- *Reactive toward:* amino groups and NHS esters/Sulfo-NHS esters or EDC-activated carboxyl groups

Ordering Information

Product #	Description	Pkg Size	U.S. Price
22299ZZ	TFCS (N-[(ε-Trifluoroacetyl)caproyloxy]succinimide ester)	50 mg	\$ 39

Special Product

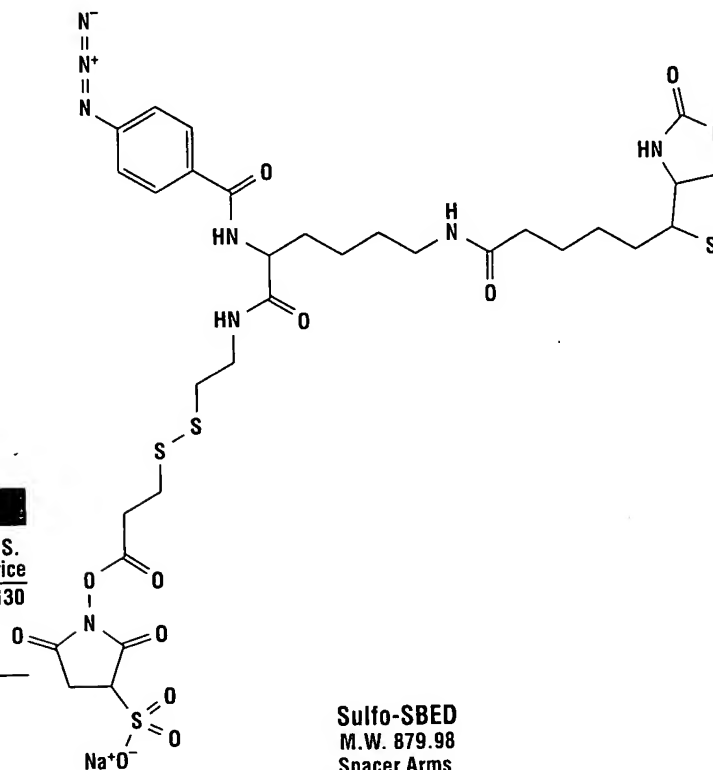


Features/Benefits:

- Amine group-specific reactivity
- Nonspecific photoreactivity
- A biotin handle; thiol-cleavable
- Soluble in DMF, DMSO and MeOH
- Moisture and light sensitive
- *Reactive groups:* phenyl azide, Sulfo-NHS ester and biotin
- *Reactive toward:* amino groups and avidin/streptavidin/NeutrAvidin™ Biotin-Binding Protein
- Literature reference #'s 49, 50, 87, 91 (page 214)

Ordering Information

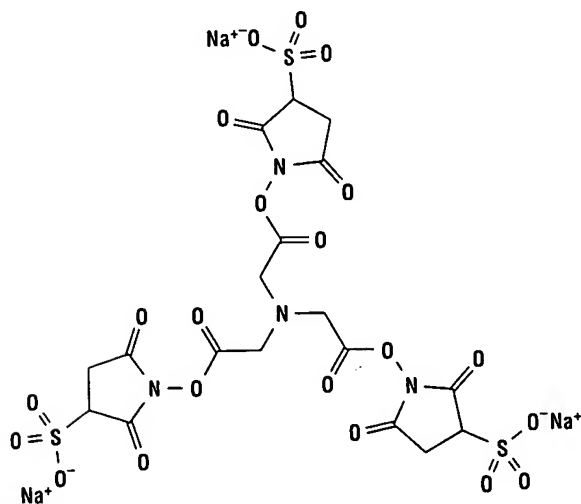
Product #	Description	Pkg. Size	U.S. Price
33033ZZ	Sulfo-SBED (Sulfosuccinimidyl [2-6-(biotinamido)-2-(p-azidobenzamido)-hexanoamido]ethyl-1,3'-dithiopropionate)	10 mg	\$130



Sulfo-SBED
M.W. 879.98
Spacer Arms
Sulfo-NHS ester 13.7 Å
Phenyl azide 9.1 Å
Biotin 19.1 Å

Sulfo-TSAT

Water-soluble, amine-reactive agent for preparing trimeric aggregates.



Sulfo-TSAT
M.W. 788.49
Spacer Arm 4.2 Å

Features/Benefits:

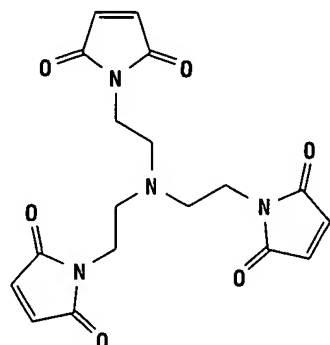
- Novel amine-reactive trifunctional cross-linking agent
- Sulfo-NHS ester couples to amines at pH 7-9 to form stable amide bonds
- Useful for the preparation of multicomponent aggregates
- Non-cleavable; water-soluble
- Literature in nearest neighbor studies
- Core molecule for the construction of dendritic polymers
- *Reactive group:* Sulfo-NHS ester
- *Reactive toward:* amino groups

Ordering Information

Product #	Description	Pkg. Size	U.S. Price
33053ZZ	Sulfo-TSAT (Tris-sulfosuccinimidyl aminotriacetate)	50 mg	\$123

TMEA

Sulfhydryl-reactive tool for preparing trimeric aggregates.



TMEA
M.W. 386.36
Spacer Arm 10.3 Å

Features/Benefits:

- Novel sulfhydryl-reactive trifunctional cross-linking agent
- Maleimides react with -SH groups at a pH of 6.5-7.5, forming stable thioether linkages
- Useful for the preparation of trimeric complexes of cysteine-containing peptides and other thiol-containing compounds
- Non-cleavable; water-insoluble
- Application in nearest neighbor studies
- Core molecule for the construction of dendritic polymers
- *Reactive group:* maleimide
- *Reactive toward:* sulfhydryl groups

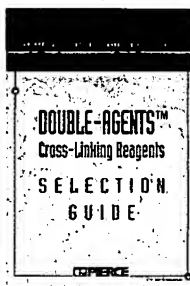
Ordering Information

Product #	Description	Pkg Size	U.S. Price
33043ZZ	TMEA (Tris-[2-maleimidoethyl]amine)	50 mg	\$109

Cross-linking Reagent Buffer Scheme

Type of Cross-linker	Buffer
Imidoester	BupH™ Borate Buffer Packs, pH 8.5 (Product # 28384ZZ) or BupH™ Carbonate-Bicarbonate Buffer Packs, pH 9.0 (Product # 28382ZZ)
Homobifunctional NHS-Ester Cross-linking Buffer	BupH™ Phosphate Buffered Saline Packs, pH 7.2 (Product # 28372ZZ) or BupH™ Modified Dulbecco's Phosphate Buffered Saline Packs, pH 7.4 (Product # 28374ZZ)
Heterobifunctional NHS-Ester Maleimide	BupH™ Phosphate Buffered Saline Packs, pH 7.2 (Product # 28372ZZ) or BupH™ Modified Dulbecco's Phosphate Buffered Saline Packs, pH 7.4 (Product # 28374ZZ)
Carbodiimide	BupH™ MES Buffered Saline, pH 4.7 (Product # 28390ZZ)
EDC with NHS or Sulfo-NHS	BupH™ Phosphate Buffered Saline Packs, pH 7.2 (Product # 28372ZZ) or BupH™ Modified Dulbecco's Phosphate Buffered Saline Packs, pH 7.4 (Product # 28374ZZ)

Double-Agents™ Cross-Linking Reagents Selection Guide



Contact Pierce, the world leader in cross-linking reagents for life science applications, for a free copy of the **Double-Agents™ Cross-Linking Reagents Selection Guide**. Written for both the novice and the expert, this guide helps you select the best reagent(s) based on your criteria and intended application. Having helped you narrow your choices, the guide provides other relevant information (molecular weight, spacer arm length, structure, etc.) to further help you select a reagent.

Ordering Information

Product #	Description
1600250	Double-Agents™ Cross-Linking Reagents Selection Guide

In a hurry to select a cross-linker? Log on to the Pierce web site at www.piercenet.com and use our online selection guide.

• 800-874-3723

Pierce Cross-linkers at a Glance

Double-Agents™ Cross-linker Product # Acronym	Reactive Toward						Cleavable By				Iodina- table		Latent Functional Group
	-NH ₂ Amines	-SH Sulf- hydryls	Carbo- hydrates	Nonselec- tive (photo- reactive)	-COOH Carboxyls	-OH Hydroxyl	Thiols	Base	Periodate	Hydroxyl- amine	Yes	No	
21509ZZ ABH			X	X								X	
22101ZZ AEDP	X				X		X					X	
22295ZZ AMAS	X	X										X	
21451ZZ ANB-NOS	X			X								X	
27720ZZ APDP		X		X			X				X		
20108ZZ APG				X								X	
21512ZZ ASBA				X	X						X		
21564ZZ BASED				X			X				X		
22331ZZ BMB		X										X	
22332ZZ BMDDB		X							X			X	
22330ZZ BMH		X										X	
22323ZZ BMOE		X										X	
22296ZZ BMPPA	X	X										X	
22297ZZ BMPH		X	X									X	
22298ZZ BMPS	X	X										X	
22336ZZ BM[PEO] ₃		X										X	
22337ZZ BM[PEO] ₄		X										X	
21600ZZ BSOCES	X							X				X	
21580ZZ BS ³	X											X	
21525ZZ DFDNB	X											X	
20663ZZ DMA	X											X	
21666ZZ DMP	X											X	
20700ZZ DMS	X											X	
21702ZZ DPDPB		X					X					X	
20593ZZ DSG	X											X	
22585ZZ DSP	X						X					X	
21555ZZ DSS	X											X	
20589ZZ DST	X								X			X	
20665ZZ DTBP	X						X					X	
22335ZZ DTME		X					X					X	
21578ZZ DTSSP	X						X					X	
22980ZZ EDC	X				X							X	
21565ZZ EGS	X									X		X	
22306ZZ EMCA	X	X										X	
22106ZZ EMCH		X	X									X	
22308ZZ EMCS	X	X										X	
22309ZZ GMBS	X	X										X	
22334ZZ HBVS		X										X	
22211ZZ KMUA	X	X										X	
22111ZZ KMHU		X	X									X	
22362ZZ LC-SMCC	X	X										X	
21651ZZ LC-SPDP	X	X					X					X	
22311ZZ MBS	X	X										X	

Internet: <http://www.piercenet.com>

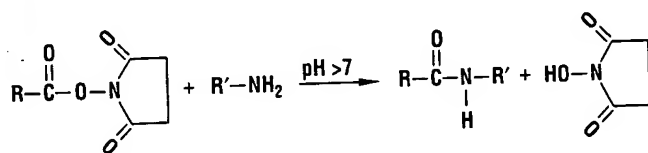
3-3
Protein Chemistry
209

*Trifunctional cross-linking reagent; binds to Avidin, Streptavidin and NeutrAvidin™ Biotin-Binding Protein. **Trifunctional cross-linking agent.

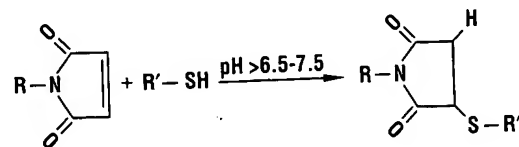
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Active Group Reaction Schemes

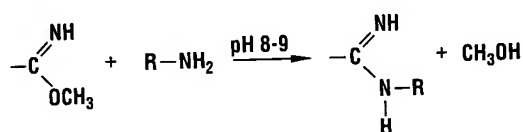
NHS-Ester Reaction Scheme



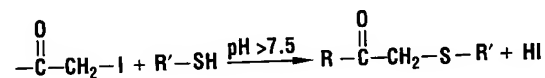
Maleimide Reaction Scheme



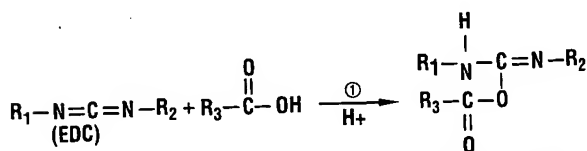
Imidoester Reaction Scheme



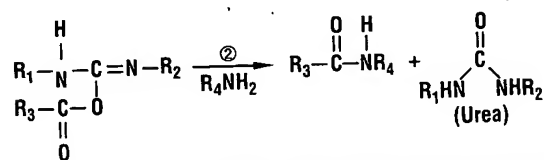
Active Halogen Reaction Scheme



EDC Coupling Reaction Scheme

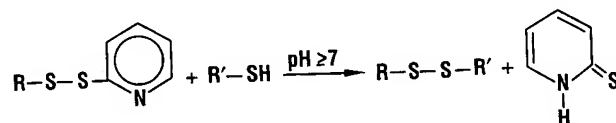


EDC reacts with carboxylic acid group and activates the carboxyl group, allowing it to be coupled to the amino group (R_4NH_2) in the reaction mixture.



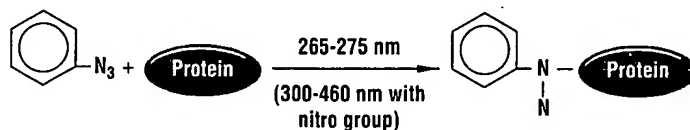
EDC is released as a soluble urea derivative after displacement by the nucleophile, R_4NH_2 .

Pyridyl Disulfide Reaction Scheme

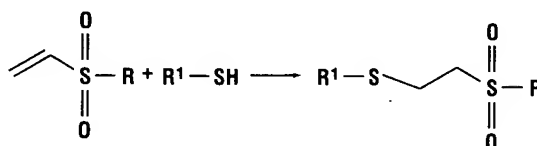


Active Group Reaction Schemes

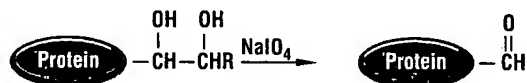
Azidophenyl Photolysis



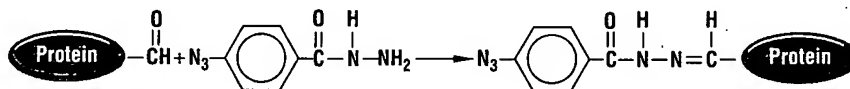
Vinyl-Sulfone Reaction Scheme



Hydrazide Reaction Scheme



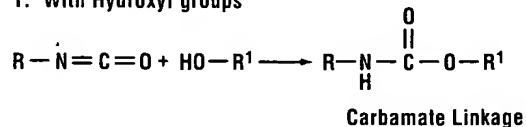
The oxidation of a Protein Carbohydrate (*cis*-diol) to an aldehyde.



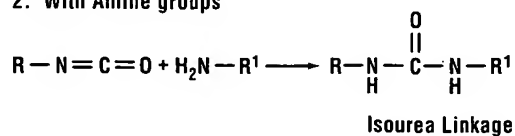
ABH, or Azidobenzoyl Hydrazide, reacts with the aldehyde on the protein to form an arylazide activated protein.

Isocyanate Reaction Schemes

1. With Hydroxyl groups



2. With Amine groups

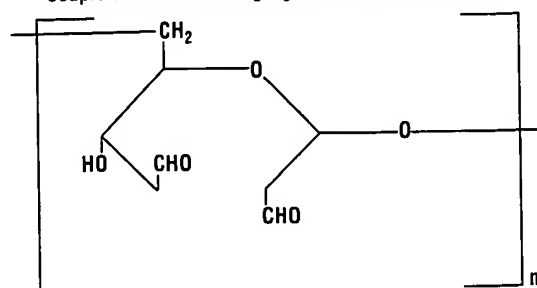


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Activated Dextran Coupling Kits

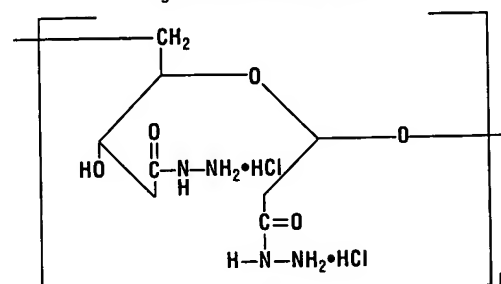
Novel water-soluble reagents capable of multimeric aggregate formation.

Couple amine-containing ligands and biomolecules



Aldehyde-Activated Dextran
Average M.W. (dextran) 40 kD

Couple aldehyde- or carboxyl-containing ligands and biomolecules



Hydrazide-Activated Dextran
Average M.W. (dextran) 40 kD

Potential Applications:

- Coupling both high and low molecular weight biomolecules to a single soluble matrix
- Serving as a water-soluble scavenger of proteases from cell preparations
- Isolating a target antigen from solution containing dextran-coupled antibodies
- Isolating a target antibody from solution via a specific ligand
- Stabilizing enzymes for research or process-level applications
- Forming antibody-enzyme conjugates for use in immunoassay development
- Preparing immunoconjugates
- Acting as a poly-functional cross-linking agent to capture multicomponent systems
- Improving the solubility properties of isolated membrane proteins

Reference

1. Mallia, A.K. and Vigna, R.A. (1998). Nonimmunogenic-activated carriers: aldehyde-activated and hydrazide-activated dextran. *Previews* 1(4), 16-17.

Ordering Information

Product #	Description	Pkg Size	U.S. Price
20890ZZ	Aldehyde-Activated Dextran Coupling Kit Includes: Aldehyde-Activated Dextran* [CHO Loading: ~300 moles/mole of dextran] Sodium Cyanoborohydride BupH™ Phosphate Buffered Saline	Kit 5 x 5 mg 190 mg 1 pack	\$ 97
20900ZZ	Hydrazide-Activated Dextran Coupling Kit Includes: Hydrazide-Activated Dextran* [Hydrazide Loading: 1-2 μmoles hydrazide/mg dextran] Sodium Cyanoborohydride BupH™ Phosphate Buffered Saline	Kit 5 x 5 mg 190 mg 1 pack	\$127

*The average molecular weight of dextran used in these preparations is 40 kD.

Compatible Products

22980ZZ	EDC (1-Ethyl-3-[3-Dimethylaminopropyl] carbodiimide Hydrochloride)	5 gm	\$ 61
22981ZZ	EDC	25 gm	\$198
28372ZZ	BupH™ Phosphate Buffered Saline Packs Each pack yields 500 ml of 0.1 M phosphate, 0.15 M NaCl, pH 7.2 when dissolved in 500 ml of distilled water.	40 packs	\$ 83
44892ZZ	AminoLink® Reductant (Sodium Cyanoborohydride)	2 gm	\$ 25

Pierce Cross-linking Reagent Literature References

1. O'Shannessy, D.J. and Quarles, R.H. (1985). Specific conjugation reactions of the oligosaccharide moieties of immunoglobulins. *J. Applied Biochem.* **7**, 347-355.
2. Krieg, U.C., Walter, P. and Johnson, A.E. (1986). Photocross-linking of the signal sequence of nascent prolactin to the 54-kilodalton polypeptide of the signal recognition particle. *Proc. Natl. Acad. Sci. USA* **83**, 8604-8606.
3. Traut, R.R., et al. (1989). *Protein Function, A Practical Approach*. Oxford: IRL Press, p. 101.
4. Sgro, J., Jacrot, B. and Chroboczek, J. (1986). *Eur. J. Biochem.* **154**, 69-76.
5. Hermanson, G.T. (1996). *Bioconjugate Techniques*, San Diego: Academic Press, pp. 284, 416.
6. Hermanson, G.T. (1996). *Bioconjugate Techniques*, San Diego: Academic Press, pp. 214, 416.
7. Chen, L.L., Rosa, J.J., Turner, S. and Pepinsky, R.B. (1991). Production of multi-meric forms of CD4 through a sugar-based cross-linking strategy. *J. Biol. Chem.* **266**(27), 18237-18243.
8. Bouizar, Z., Fouchereau-Person, M., Taboulet, J., Moukhtar, M.S. and Milhaud, G. (1986). Purification and characterization of calcitonin receptors in rat kidney membranes by covalent cross-linking techniques. *Eur. J. Biochem.* **155**, 141-147.
9. Knoller, S., Shpungin, S. and Pick, E. (1991). The membrane-associated component of the amphiphile-activated, cytosol-dependent superoxide-forming NADPH oxidase of macrophages is identical to cytochrome b559. *J. Biol. Chem.* **266**, 2795-2804.
10. Kornblatt, J.A. and Lake, D.F. (1980). Cross-linking of cytochrome oxidase subunits with difluorodinitrobenzene. *Can. J. Biochem.* **58**, 219-224.
11. Hartman, F.C. and Wold, F. (1967). Cross-linking of bovine pancreatic ribonuclease A with dimethyl adipimidate. *Biochem.* **6**(8), 2439-2448.
12. Schneider, C., Newman, R.A., Sutherland, D.R., Asser, U. and Greaves, M.F. (1982). A one-step purification of membrane proteins using a high efficiency immunomatrix. *J. Biol. Chem.* **257**(18), 10766-10769.
13. Wang, D. and Moore, S. (1977). Polyspermine-ribonuclease prepared by cross-linkage with dimethyl suberimidate. *Biochem.* **16**(13), 2937-2941.
14. Chen, L.L., Frankel, A.D., Harder, J.L., Fawell, S., Barsoun, J. and Pepinsky, B. (1995). Increased cellular uptake of the human immunodeficiency virus-1 tat protein after modification with biotin. *Anal. Biochem.* **227**, 168-175.
15. Waugh, S.M., DiBella, E.E. and Pilch, P.F. (1989). Isolation of a proteolytically derived domain of the insulin receptor containing the major site of cross-linking/binding. *Biochemistry* **28**, 3448-3455. (EGS example)
16. Joshi, S. and Burrows, R. (1990). AT synthase complex from bovine heart mitochondria. *J. Biol. Chem.* **265**, 14518-14525.
17. Cox, G.W., Mattieson, B.J., Giardina, S.L. and Varesio, L. (1990). Characterization of IL-2 receptor expression and function on murine macrophages. *J. Immunol.* **145**, 1719-1726.
18. Farries, T.C. and Atkinson, J.P. (1989). Biosynthesis of properdin. *J. Immunol.* **142**, 842-847.
19. Shivdasani, R.A. and Thomas, D.W. (1988). Molecular associations of IA antigens after T-B cell interactions. *J. Immunol.* **141**, 1252-1260.
20. Jung, S.M. and Moroi, M. (1983). Cross-linking of platelet glycoprotein Ib by *N*-succinimidyl(4-azidophenyl)dithio)propionate and 3,3'-dithiobis(sulfosuccinimidyl propionate). *Biochem. Biophys. Acta* **761**, 152-162.
21. Taniuchi, M., Clark, H.B. and Johnson, Jr., E.M. (1986). Induction of nerve growth factor receptor in Schwann cells after axotomy. *Proc. Natl. Acad. Sci. USA* **83**, 4094-4098.
22. Millar, J.B. and Rozengur, E. (1990). Chronic desensitization to bombesin by progressive down-regulation of bombesin receptors in Swiss 3T3 cells. *J. Biol. Chem.* **265**, 12052-12058.
23. Fujiwara, K., et al. (1988). Sandwich enzyme immunoassay of tumor-associated antigen sialosylated Lewis (x) using β -D-galactosidase coupled to a monoclonal antibody of IgM isotype. *J. Immunol. Meth.* **112**, 77-83.
24. Ballmer-Hofer, K., Schlup, V., Burn, P. and Burger, M.M. (1982). Isolation of *in situ* cross-linked ligand-receptor complexes using an anticross-linker specific antibody. *Anal. Biochem.* **126**, 246-250.
25. Kitagawa, T. and Aikawa, T. (1976). Enzyme coupled immunoassay of insulin using a novel coupling reagent. *J. Biochem. (Tokyo)* **79**, 233-236.
26. Chamow, S.M., Kogan, T.P., Peers, D.H., Hastings, R.C., Byrn, R.A. and Askenaszi, A. (1992). Conjugation of soluble CD4 without loss of biological activity via a novel carbohydrate-directed cross-linking reagent. *J. Biol. Chem.* **267**(22), 15916-15922.
27. van der Horst, G.T.J., Mancini, G.M.S., Brossmer, R., Rose, U. and Verheijen, F.W. (1990). Photoaffinity labeling of a bacterial sialidase with an aryl azide derivative of sialic acid. *J. Biol. Chem.* **265**(19), 10801-10804. (NHS-ASA example)
28. Greenfield, R.S., Kaneko, T., Daues, A., Edson, M.A., Fitzgerald, A., Olech, L.J., Grattan, J.A., Spitalny, G.L. and Braslawsky, G.R. (1990). Evaluation *in vitro* of adriamycin immunoconjugates synthesized using an acid sensitive hydrazone linker. *Cancer Res.* **50**, 6600-6607.
29. Lewis, R.V., Roberts, M.F., Dennis, E.A. and Allison, W.S. (1977). Photoactivated heterobifunctional cross-linking reagents which demonstrate the aggregation state of phospholipase A2. *Biochemistry* **16**, 5650-5654. (ANB-NOS example)
30. Wood, C.L. and O'Dorisio, M.S. (1985). Covalent cross-linking of vasoactive intestinal polypeptide to its receptors on intact human lymphoblasts. *J. Biol. Chem.* **260**, 1243-1247. (HSAB example)
31. Chattopadhyay, A., James, H.I. and Fair, D.S. (1992). Molecular recognition sites on factor Xa which participate in the prothrombinase complex. *J. Biol. Chem.* **267**, 12323-12329.
32. Kitagawa, T., Shimozono, T., Aikawa, T., Yoshida, T. and Nishimura, H. (1981). Preparation and characterization of heterobifunctional cross-linking reagents for protein modifications. *Chem. Pharm. Bull.* **29**(4), 1130-1135.
33. Cumber, A.J., Forrester, J.A., Foxwell, B.M.J., Ross, W.C.J. and Thorpe, P.W. (1985). Preparation of antibody-toxin conjugates. *Meth. Enzymol.*, New York: Academic Press, **112**, pp. 207-225.
34. Hermanson, G.T. (1996). *Bioconjugate Techniques*, San Diego: Academic Press, pp. 542, 553, 568.
35. Uto, I., Ishimatsu, T., Hirayama, H., Ueda, S., Tsuruta, J. and Kambara, T. (1991). Determination of urinary Tamm-Horsfall protein by ELISA using a maleimide method for enzyme-antibody conjugation. *J. Immunol. Meth.* **138**, 87-94.
36. Iwai, K., Fukuoka, S.-I., Fushiki, T., Kido, K., Sengoku, Y. and Semba, T. (1988). Preparation of a verifiable peptide-protein immunogen: direction-controlled conjugation of a synthetic fragment of the monitor peptide with myoglobin and application for sequence analysis. *Anal. Biochem.* **171**, 277-282.
37. Ghetie, V., et al. (1990). *Bioconjugate Chem.* **1**, 24-31.
38. Carlsson, J., Drevin, H. and Axen, R. (1978). Protein thiolation and reversible protein-protein conjugation. *N*-succinimidyl-3-(2-pyridyldithio)propionate, a new heterobifunctional reagent. *Biochem. J.* **173**, 723-737.
39. Zarling, D.A., Watson, A. and Bach, F.H. (1980). Mapping of lymphocyte surface polypeptide antigens by chemical cross-linking with BSOCOES. *J. Immunol.* **124**, 913-920.
40. Park, L.S., Friend, D., Gillis, S. and Urdal, D.L. (1986). Characterization of the cell surface receptor for a multi-lineage colony-stimulating factor (CSF-2). *J. Biol. Chem.* **261**, 205-210. (DST example)
41. Browning, J. and Ribolini, A. (1989). Studies on the differing effects of tumor necrosis factor and lymphotoxin on the growth of several human tumor lines. *J. Immunol.* **143**, 1859-1867. (EGS example)
42. Myers, D.E., Uckun, F.M., Swaim, S.E. and Vallera, D.A. (1989). The effects of aromatic and aliphatic maleimide cross-linkers and anti-CD-5 ricin immunotoxins. *J. Immunol. Meth.* **121**, 129-142. (MBS example)

Cross-linking Reagents

Pierce Cross-linking Reagent Literature References *continued*

43. Kitagawa, T. (1981). Enzyme labeling with *N*-hydroxysuccinimidyl ester of maleimide. *Enzyme Immunoassay*, Tokyo/New York, Igaku-Shoin pp. 81-89.
44. Hermanson, G.T. (1996). *Bioconjugate Techniques*, San Diego: Academic Press, pp. 266-268.
45. Hermanson, G.T. (1996). *Bioconjugate Techniques*, San Diego: Academic Press, pp. 239-242.
46. Samoszuk, M.K., Petersen, A., Lo-Hsueh, M. and Rietveld, C. (1989). A peroxide-generating immunoconjugate directed to eosinophil peroxidase is cytotoxic to Hodgkin's disease cells *in vitro*. *Antibody, Immunoconjugates and Radiopharmaceuticals* **2**(1), 37-45.
47. Teale, J.M. and Kearney, J.R. (1986). Clonotypic analysis of the fetal B cell repertoire: Evidence for an early and predominant expression of idiotypes associated with the VH 36-60 family. *J. Mol. Cell. Immunol.* **2**, 283-292.
48. Hermanson, G.T. (1996). *Bioconjugate Techniques*, San Diego: Academic Press, pp. 232-235.
49. Hermanson, G.T. (1996). *Bioconjugate Techniques*, San Diego: Academic Press, pp. 289, 291, 375.
50. *Previews*, May/June 1994, p. 1.
51. Lala, A., Sojar, H.T. and DeNardin, E. (1996). Quantitative approach for detection and characterization of formyl peptide receptor in solution using a photo affinity ligand. *Peptide Res.* **9**(2), 58-60.
52. Bieniarz, C., Husain, M., Barnes, G., King, C.A. and Welch, C.J. (1996). Extended length heterofunctional coupling agents for protein conjugations. *Bioconjugate Chem.* **7**, 88-95.
53. Vanin, E.F. and Ji, T.H. (1981). Synthesis and application of cleavable photoactivable heterobifunctional reagents. *Biochemistry* **20**, 6754-6760.
54. Thevenin, B., et al. (1991). A novel reagent for functionally directed site-specific fluorescent labeling of proteins. *Biophys. J.* **59**, 358a.
55. Eager, J.E. and Savage, W.E. (1963). Photolysis and photo-oxidation of amino acids and peptides - VI. A study of the initiation of disulfide interchange by light irradiation. *Photochem. Photobiol.* **2**, 25-37.
56. Schnaar, R.L., Langer, B.G. and Brandley, B.K. (1985). Reversible covalent immobilization of ligands and proteins on polyacrylamide gels. *Anal. Biochem.* **151**, 268-281.
57. May, J.M. (1989). Selective labeling of the erythrocyte hexose carrier with a maleimide derivative of glucosamine: Relationship of an exofacial sulfhydryl to carrier conformation and structure. *Biochemistry* **28**, 1718-1725.
58. Sayre, L.M., Larson, D.L., Takemori, A.E. and Portoghesi, P.S. (1984). Design and synthesis of naltrexone-derived affinity labels with nonequilibrium opoid agonist and antagonist activities. Evidence for the existence of different mu receptor subtypes in different tissues. *J. Med. Chem.* **27**(10), 1325-35.
59. Stalteri, M.A. and Mather, S.J. (1995). A cross-linked monoclonal antibody fragment for improved tumor targeting. *Bioconjugate Chem.* **6**, 179-186.
60. Yi, F., Denker, B.M. and Neer, E.J. (1991). Structural and functional studies of cross-linked G0 protein subunits. *J. Biol. Chem.* **266**(6), 3900-3906.
61. Hermanson, G.T. (1996). *Bioconjugate Techniques*, San Diego, Academic Press, pp.114-116, 294. (periodate cleavage of vic-diols)
62. O'Sullivan, M., et al. (1979). Comparison of two methods of preparing enzyme-antibody conjugates: Application of these conjugates to enzyme immunoassay. *Anal. Biochem.* **100**, 100-108.
63. Rich, D., Geselchen, P.D., Tong, D., Cheung, A. and Buckner, C.K. (1975). Alkylating derivatives of amino acids and peptides. Synthesis of *N*-maleoylamino acids, [1-(*N*-maleoyl)glycyl]cysteinylloxycytosine. Effects on vasopressin-stimulated water loss from isolated toad bladder. *J. Med. Chem.* **18**, 1004-1010.
64. Moroder, L., et al. (1983). Immunoassays of peptide hormones and their chemical aspects. *Biopolymers* **22**, 481-486.
65. Han, J.C. and Han, G.Y. (1994). A procedure for quantitative determination of tris(2-carboxyethyl)phosphine, an odorless reducing agent more stable and effective than dithiothreitol. *Anal. Biochem.* **220**, 5-10.
66. Griffith, D.G., Partis, M.D., Sharp, R.N. and Beechey, R.B. (1981). *N*-Polymethylenecarboxymaleimides. A new class of probes for membrane sulfhydryl groups. *FEBS Lett.* **134**, 261-263.
67. Trail, P.A., et al. (1993). Cure of xenografted human carcinomas by BR96-doxorubicin immunoconjugates. *Science* **261**, 212-215.
68. Peeters, J.M., Hazendonk, T.G., Beuvery, E.C. and Tesser, G.I. (1989). Comparison of four bifunctional reagents for coupling peptides to protein and the effect of these three moieties on the immunogenicity of the conjugates. *J. Immunol. Methods* **120**, 133-143.
69. Hermanson, G.T. (1996). *Bioconjugate Techniques*, San Diego: Academic Press, pp. 243-245. (EMCS use can be modeled after GMBS.)
70. Yoshitake, S., Yamada, Y., Ishikawa, E. and Masseyeff, R. (1979). Conjugation of glucose oxidase from *Aspergillus niger* and rabbit antibodies using *N*-(4-carboxycyclohexylmethyl)-maleimide. *Eur. J. Biochem.* **101**, 395-399.
71. Annunziato, M.E., Patel, U.S., Ranade, M. and Palumbo, P.S. (1993). *p*-Maleimidophenyl isocyanate: A novel heterobifunctional linker for hydroxyl to thiol coupling. *Bioconjugate Chem.* **4**, 212-218.
72. Duncan, R.J.S., Weston, P.D. and Wigglesworth, R. (1983). A new reagent which may be used to introduce sulfhydryl groups into protein, and its use in the preparation of conjugates for immunoassay. *Anal. Biochem.* **132**, 68-73.
73. Inman, J.K., Highet, P.F., Kolodny, N. and Robey, F.A. (1991). Synthesis of *N*-alpha-(tert-butoxycarbonyl)-*N*-epsilon-[*N*-(bromoacetyl)-beta-alanyl]-L-lysine: Its use in peptide synthesis for placing a bromoacetyl cross-linking function at any desired sequence position. *Bioconjugate Chem.* **2**, 458-463.
74. Thorpe, P.E., et al. (1984). Blockade of the galactose-binding sites of ricin by its linkage to antibody. Specific cytotoxic effects of the conjugates. *Eur. J. Biochem.* **140**, 63-71.
75. Rector, E.S., Schwenk, R.J., Tse, K.S. and Sehon, A.H. (1978). A method for the preparation of protein-protein conjugates of predetermined composition. *J. Immunol. Methods* **24**, 321-336.
76. Zera, J.J., et al. (1991). A carbohydrate-directed heterobifunctional cross-linking reagent for the synthesis of immunoconjugates. *Anal. Biochem.* **194**, 156-162.
77. Friden, P.M., et al. (1993). Blood-brain barrier penetration and *in-vivo* activity of an NGF conjugate. *Science* **259**, 373-377.
78. Masri, M.S. and Friedman, M. (1988). Protein reactions with methyl and ethyl vinyl sulfones. *J. Protein Chem.* **7**, 49-54.
79. Morpurgo, M., et al. (1996). Preparation and characterization of poly(ethylene glycol)vinyl sulfone. *Bioconjugate Chem.* **7**, 363-368.
80. Ishii, Y. and Lehrer, S.S. (1986). Effects of the state of the succinimido-ring on the fluorescence and structural properties of pyrene maleimide-labeled alpha alpha-tropomyosin. *Biophys. J.* **50**, 75-89.
81. Fujii, N., Akaji, K., Hayashi, Y. and Yajima, H. (1985). Studies on peptides. CXXV. 3-(3-*p*-methoxybenzylthiopropionyl)-thiazolidine-2-thione and its analogs as reagents for the introduction of the mercapto group into peptides and proteins. *Chem. Pharm. Bull.* **33**, 362-367.
82. Pandurangi, R.S., Lusiak, R., Desai, S. and Kuntz, R.R. (1998). Chemistry of bifunctional photoprobes: Part 4: Synthesis of the chromogenic, cleavable, water soluble and heterobifunctional (*N*-Methyl amino perfluoroaryl azide benzamido)-ethyl-1,3-dithiopropionyl sulfosuccinimide: An efficient protein cross-linking agent. *J. Chem. Soc. Chem. Commun.* Submitted for publication.
83. Pandurangi, R.S., Karra, R.S. and Volkert, W.A. (1997). Recent trends in the evaluation of the photochemical insertion characteristics of heterobifunctional perfluoroaryl azide chelating agents: Biochemical implication in nuclear medicine. *Photochem. Photobiol.* **65**(2), 208-221.
84. Chrisey, L.A., Lee, G.U. and O'Ferrall, C.E. (1996). Covalent attachment of synthetic DNA to self-assembled monolayer films. *Nucleic Acids Research* **24**(15), 3031-3039.
85. Rajur, S.B., Roth, C.M., Morgan, J.R. and Yarmush, M.L. (1997). Covalent protein-oligonucleotide conjugates for efficient delivery of antisense molecules. *Bioconjugate Chem.* **8**, 935-940.
86. Wang, D., et al. (1997). Generation and characterization of an anti-CD19 single-chain Fv immunotoxin composed of C-terminal disulfide-linked dgRTA. *Bioconjugate Chem.* **8**, 878-884.
87. Liver, D., et al. (1998). *Helicobacter pylori* adhesion binding fucosylated histo-blood group antigens revealed by retagging. *Science* **279**, 373-376.
88. Grabarek, Z. and Gergely, J. (1990). Zero-length cross-linking procedure with the use of active esters. *Anal. Biochem.* **185**(1), 131-135.
89. Kuipers, W.H., Bos, E.S., Kaspersen, F.M., Veenman, G.H. and van Boeckel, C.A. (1993). Specific recognition of antibody-oligonucleotide conjugates by radiolabeled antisense nucleotides: a novel approach for two step radio-immunotherapy of cancer. *Bioconjugate Chem.* **4**(1), 94-102.
90. Nakagami, S., Matsunaga, H., Oka, N. and Yamane, A. (1991). Preparation of enzyme-conjugated DNA probe and application to the universal probe system. *Anal. Biochem.* **198**(1), 75-79.
91. Geselowitz, D.A. and Neumann, R.D. (1995). Quantitation of triple-helix formation using a photo-cross-linkable arylazide/biotin/oligonucleotide conjugate. *Bioconjugate Chem.* **6**(4), 502-506.